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STRUCTURE FILE UPDATES: 7 MAR 2003 HIGHEST RN 497212-14-3 DICTIONARY FILE UPDATES: 7 MAR 2003 HIGHEST RN 497212-14-3

TSCA INFORMATION NOW CURRENT THROUGH MAY 20, 2002

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FILE 'CA' ENTERED AT 07:44:57 ON 09 MAR 2003 COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)

にょに 89 L1 AND CONOTOXIN

ユロ ANSWER 1 OF 89 CA COPYRIGHT 2003 ACS

Identification of the nicotinic receptor subtypes expressed on dopaminergic terminals in the rat striatum PY

ロロ ANSWER 2 OF 89 CA COPYRIGHT 2003 ACS

Declines in different .beta.2* nicotinic receptor populations in monkey striatum after nigrostriatal damage PY 2002

ゴロ ANSWER 3 OF 89 CA COPYRIGHT 2003 ACS

.alpha.- ***Conotoxin *** IC from Conus geographus, a Novel Peptide Antagonist of Nicotinic Acetylcholine Receptors PY 2002

ゴロ ANSWER 4 OF 89 CA COPYRIGHT 2003 ACS

The synthesis and structure of an n-terminal dodecanoic acid conjugate of .alpha. ***conotoxin*** M Py

ANSWER 5 OF 89 CA COPYRIGHT 2003 ACS

Characterization of [125] ppibatidine binding and nicotinic agonist-mediated 86Rb+ efflux in interpeduncular nucleus and inferior colliculus of .beta.2 null mutant mice PY 2002

ANSWER 6 OF 89 CA COPYRIGHT 2003 ACS

ユロ Toxins 'R' Us: more pharmacological tools from nature's superstore PY 2002

ゴロ ANSWER 7 OF 89 CA COPYRIGHT 2003 ACS

peptides with analgesic properties PY

L4 ANSWER 8 OF 89 CA COPYRIGHT 2003 ACS TI Differential nicotinic receptor expression in monke Differential nicotinic receptor expression in monkey basal ganglia: Effects of nigrostriatal damage PY

L4 ANSWER 9 OF 89 CA COPYRIGHT 2003 ACS TI Methyllycaconitine is a potent antagonist of .alpha PY 2002 Methyllycaconitine is a potent antagonist of .alpha.- ***conotoxin*** -MII-sensitive presynaptic nicotinic acetylcholine receptors in rat striatum

L4 ANSWER 10 OF 89 CA COPYRIGHT 2003 ACS

TI Novel conditions for use in the therapeutic regulation of ion channel function PY 2002

L4 ANSWER 11 OF 89 CA COPYRIGHT 2003 ACS TI A novel choline-sensitive nicotinic receptor subtype

A novel choline-sensitive nicotinic receptor subtype that mediates enhanced GABA release in the chick ventral lateral geniculate nucleus PY

subtypes PY L4 ANSWER 12 OF 89 CA COPYRIGHT 2003 ACS TI 5-lodo-A-85380 binds to .alpha.- ***conntoxin*** 5-lodo-A-85380 binds to .alpha.- ***conotoxin*** MII-sensitive nicotinic acetylcholine receptors (nAChRs) as well as .alpha.4.beta.2* btypes PY 2002

L4 ANSWER 13 OF 89 CA COPYRIGHT 2003 ACS
T1 Involvement of the .alpha.3 subunit in central nicotinic binding populations PY

L4 ANSWER 14 OF 89 CA COPYRIGHT 2003 ACS TI Solution conformation of .alpha.- ***Conotoxin*** nicotinic acetylcholine receptor PY Solution conformation of .alpha.- ***Conotoxin*** El, a neuromuscular toxin specific for the .alpha.1/.delta. subunit interface of Torpedo

ロロ ANSWER 15 OF 89 CA COPYRIGHT 2003 ACS

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ANSWER 16 OF 89 CA COPYRIGHT 2003 ACS

Protein sequences of synthetic Conus .alpha -conotoxins and the therapeutic uses thereof as neuromuscular blocking agent PY 2002

> TI New members of the .mu. - ***conotoxin*** family for use in the treatment of disease associated with sodium channel function and cDNAs encoding them PY 2002 2002

ANSWER 17 OF 89 CA COPYRIGHT 2003 ACS

conotoxin MII sites PY 2002 Loss of nicotinic receptors in monkey striatum after 1-methyl-4-phenyl- 1,2,3,6-tetrahydropyridine treatment is due to a decline in .alpha.

ANSWER 18 OF 89 CA COPYRIGHT 2003 ACS

Two new classes of conopeptides inhibit the .alpha. 1-adrenoceptor and noradrenaline transporter PY 2 2 2 3

ANSWER 19 OF 89 CA COPYRIGHT 2003 ACS

Vulnerability of 1251-alpha.- ***conotoxin*** MII binding sites to nigrostriatal damage in monkey PY 200

ANSWER 20 OF 89 CA COPYRIGHT 2003 ACS

ロロロ Mechanisms for evolving hypervariability: the case of conopeptides PY

⊒ ⊑ ANSWER 21 OF 89 CA COPYRIGHT 2003 ACS

An efficient synthetic scheme for natural .alpha.-conotoxins and their analogues PY

L4 ANSWER 22 OF 89 CA COPYRIGHT 2003 ACS

TI An .alpha.4.beta.4 nicotinic receptor subtype is present in chick retina: identification, characterization and pharmacological comparison with the transfected .alpha.4.beta.4 and .alpha.6.beta.4 subtypes PY 2001

⊒ ⊑ ANSWER 23 OF 89 CA COPYRIGHT 2003 ACS

Structure-Activity Relationships in a Peptidic .alpha.7 Nicotinic Acetylcholine Receptor Antagonist PY 2000

L4 ANSWER 24 OF 89 CA COPYRIGHT 2003 ACS
TI Protein and cDNA sequences of Conus conotoxins and therapeutic uses thereof PY

L4 ANSWER 25 OF 89 CA COPYRIGHT 2003 ACS TI Alpha-conotoxins and nucleic acids encoding them PY 2000 2001 2002

L4 ANSWER 26 OF 89 CA COPYRIGHT 2003 ACS

2000 2000 2001 2001 TI Protein and cDNA sequences of Conus .alpha.-conotoxins and the therapeutic uses thereof as neuromuscular blocking agent PY 200

ANSWER 27 OF 89 CA COPYRIGHT 2003 ACS 125I-alpha - ***conotoxin*** MII identifies a novel nicotinic acetylcholine receptor population in mouse brain ₽ 2000

ANSWER 28 OF 89 CA COPYRIGHT 2003 ACS

ゴロ .beta.3 Subunit is present in different nicotinic receptor subtypes in chick retina PY

L4 ANSWER 29 OF 89 CA COPYRIGHT 2003 ACS TI Conus peptides: novel probes for nicotinic acetylch

Conus peptides: novel probes for nicotinic acetylcholine receptor structure and function PY

rat striatal synaptosomes PY L4 ANSWER 30 OF 89 CA COPYRIGHT 2003 ACS
TI UB-166: A novel nicotinic agonist with subtype selectivity implicates the .alpha.4.beta.2* subtype in the modulation of dopamine release from

⊒ ⋤ ANSWER 31 OF 89 CA COPYRIGHT 2003 ACS

Leu10 of .alpha.- ***conotoxin*** PnlB confers potency for neuronal nicotinic responses in bovine chromaffin cells PY 2000

L4 ANSWER 32 OF 89 CA COPYRIGHT 2003 ACS
TI ..ho.-***Conotoxin*** peptides with alpha 1-adrenoceptor antagonist activity, nucleic acids encoding them, antibodies, and therapeutic uses PY 2000 2000 2000 2001 2002

4 ANSWER 33 OF 89 CA COPYRIGHT 2003 ACS

Pairwise interactions between neuronal alpha.7 acetylcholine receptors and alpha.- ***conotoxin*** PnIB PY

ロロ ANSWER 34 OF 89 CA COPYRIGHT 2003 ACS

peptides PY 2000

L4 ANSWER 35 OF 89 CA COPYRIGHT 2003 ACS TI Pharmacological characterization of the response of the company of the response of the company of

Pharmacological characterization of the response of the leech pharynx to acetylcholine PY

L4 ANSWER 36 OF 89 CA COPYRIGHT 2003 ACS

Single amino acid substitutions in .alpha.- ***conotoxin***
etylcholine receptor PY 1999 PnIA shift selectivity for subtypes of the mammalian neuronal nicotinic

L4 ANSWER 37 OF 89 CA COPYRIGHT 2003 ACS TI Conopeptides from Conus striatus and Conus textile

Conopeptides from Conus striatus and Conus textile by cDNA cloning PY 1999

ユロ ANSWER 38 OF 89 CA COPYRIGHT 2003 ACS

Single-Residue Alteration in .alpha.- ***Conotoxin*** PnIA Switches Its nAChR Subtype Selectivity PY 1999

ゴロ ANSWER 39 OF 89 CA COPYRIGHT 2003 ACS

Aromatic substitutions in .alpha.- ***conotoxin*** Iml. Synthesis of iodinated photoactivatable derivative PY 1999

ANSWER 40 OF 89 CA COPYRIGHT 2003 ACS

ユロ Three-dimensional structure of .alpha.- ***conotoxin*** El determined by 1H NMR spectroscopy PY

L4 ANSWER 41 OF 89 CA COPYRIGHT 2003 ACS TI Minimal conformation of the .alpha. ***conotoxin* NMR and binding studies PY Minimal conformation of the .alpha. ***conotoxin*** Iml for the .alpha.7 neuronal nicotinic acetylcholine receptor recognition: correlated CD,

ANSWER 42 OF 89 CA COPYRIGHT 2003 ACS

ゴロ Functional .alpha.6-containing nicotinic receptors are present in chick retina PY

ANSWER 43 OF 89 CA COPYRIGHT 2003 ACS

ュロ Pairwise interactions between neuronal .alpha.7 acetylcholine receptors and .alpha.- ***conotoxin*** m PY 1999

ANSWER 44 OF 89 CA COPYRIGHT 2003 ACS

Cloning and sequencing of alpha- ***conotoxin*** sequences from Conus textile venom duct PY

ANSWER 45 OF 89 CA COPYRIGHT 2003 ACS

Solution structure of .alpha.- ***conotoxin*** Iml determined by two-dimensional NMR spectroscopy PY

ANSWER 46 OF 89 CA COPYRIGHT 2003 ACS

ゴロ Solution Structure of .alpha.- ****Conotoxin*** Iml by 1H Nuclear Magnetic Resonance PY 1999

コロ ANSWER 47 OF 89 CA COPYRIGHT 2003 ACS

Inhibition of nicotine-induced hippocampal norepinephrine release in rats by alpha-conotoxins MII and AuIB microinjected into the locus

⊒ ⊏ ANSWER 48 OF 89 CA COPYRIGHT 2003 ACS Uses of alpha- ***conotoxin*** peptides PY 19 1999 1999 2001 2002

L4 ANSWER 49 OF 89 CA COPYRIGHT 2003 ACS

phosphopeptides by electrospray, matrix-assisted laser desorption/tonization (MALDI) and atmospheric pressure MALDI mass spectrometry PY 1999 TI Identification of tyrosine sulfation in Conus pennaceus conotoxins .alpha.-PnIA and .alpha.-PnIB: further investigation of labile sulfo- and

L4 ANSWER 50 OF 89 CA COPYRIGHT 2003 ACS

T1 .alpha.- ***Conotoxin*** ImI inhibits the .alpha.-bungarotoxin- resistant nicotinic response in bovine adrenal chromaffin cells PY 1999

L4 ANSWER 51 OF 89 CA COPYRIGH1 2003 AUS

TI NMR Solution Structure of .alpha.- ***Conotoxin*** Iml and Comparison to Other Conotoxins Specific for Neuronal Nicotinic Acetylcholine

L4 ANSWER 52 OF 89 CA COPYRIGHT 2003 ACS
TI Preparation and interaction of .alpha. ***conotoxin*** peptides with neuronal nicotinic acetylcholine receptors PY 1999 1999 1999 2000

L4 ANSWER 53 OF 89 CA COPYRIGHT 2003 ACS TI NMR spatial structure of .alpha. - ***conotoxin*** | acetylcholine receptors PY 1999 Iml reveals a common scaffold in snail and snake toxins recognizing neuronal nicotinic

L4 ANSWER 54 OF 89 CA COPYRIGHT 2003 ACS
TI Accelerated chemical synthesis of peptides and small proteins PY

L4 ANSWER 55 OF 89 CA COPYRIGHT 2003 ACS

Unmasking the functions of the chromaffin cell alpha.7 nicotinic receptor by using short pulses of acetylcholine and selective blockers ş

L4 ANSWER 56 OF 89 CA COPYRIGHT 2003 ACS TI .alpha.- ***Conotoxin*** AulB selectively blocks .¢ PY 1998 AulB selectively blocks .alpha.3.beta.4 nicotinic acetylcholine receptors and nicotine-evoked norepinephrine release

L4 ANSWER 57 OF 89 CA COPYRIGHT 2003 ACS TI Three-Dimensional Solution Structure of .alpha. **

Three-Dimensional Solution Structure of .alpha - ***Conotoxin*** MII by NMR Spectroscopy: Effects of Solution Environment on Helicity PY

L4 ANSWER 58 OF 89 CA COPYRIGHT 2003 ACS

Functional determinants by which snake and cone snail toxins block the .alpha.7 neuronal nicotinic acetylcholine receptors PY 1998

L4 ANSWER 59 OF 89 CA COPYRIGHT 2003 ACS

Molecular dissection of subunit interfaces in the nicotinic acetylcholine receptor PY

L4 ANSWER 60 OF 89 CA COPYRIGHT 2003 ACS Toxic conopeptides AulA, AulB and AulC of cone snail venom active against nicotinic receptors PY 1998 1999

L4 ANSWER 61 OF 89 CA COPYRIGHT 2003 ACS

neurons PY 1998 TI Two distinct nicotinic receptors, one pharmacologically similar to the vertebrate .alpha.7-containing receptor, mediate CI currents in Aplysia

L4 ANSWER 62 OF 89 CA COPYRIGHT 2003 ACS

Ti The 1.1 ANG. Resolution Crystal Structure of [Tyr15]Epl, a Novel alpha. ***Conoloxin*** from Conus episcopatus, Solved by Direct

L4 ANSWER 63 OF 89 CA COPYRIGHT 2003 ACS

Ti .alpha.- ""Conotoxin" Epi, a novel sulfated peptide from Conus episcopatus that selectively targets neuronal nicotinic acetylcholine

L4 ANSWER 64 OF 89 CA COPYRIGHT 2003 ACS TI Identification of residues in the neuronal .alpha.7 ac

Identification of residues in the neuronal .alpha.7 acetylcholine receptor that confer selectivity for ***conotoxin*** Iml PY

L4 ANSWER 65 OF 89 CA COPYRIGHT 2003 ACS

TI Use of alpha. ***conotoxin*** MII? to treat disorders resulting from nicotine-stimulated dopamine release PY 1998 1998 1998 199

L4 ANSWER 66 OF 89 CA COPYRIGHT 2003 ACS
T1 Structural elements in .alpha.7 receptors Ŗ

L4 ANSWER 67 OF 89 CA COPYRIGHT 2003 ACS

peptides Iml and MII as cardiovascular agents PY 1998 1998 2001 1999 2001

L4 ANSWER 68 OF 89 CA COPYRIGHT 2003 ACS

slices PY TI Differential inhibition by .alpha - ***conotoxin*** - MII of the nicotinic stimulation of [3H] dopamine release from rat striatal synaptosomes and

L4 ANSWER 69 OF 89 CA COPYRIGHT 2003 ACS TI Three-Dimensional Solution Structure of .alpha.- ***Conotoxin***

Mll, an .alpha.3.beta.2 Neuronal Nicotinic Acetylcholine Receptor-Targeted

L4 ANSWER 70 OF 89 CA COPYRIGHT 2003 ACS TI Crystal Structure at 1.1 ANG. Resolution of alpha.

Crystal Structure at 1.1 .ANG. Resolution of .alpha.- ***Conotoxin*** PnlB: Comparison with .alpha.-Conotoxins PnlA and GI PY 1997

L4 ANSWER 71 OF 89 CA COPYRIGHT 2003 ACS TI Determinants of specificity for .alpha.- ***conotoxin Determinants of specificity for .alpha.- ***conotoxin***

L4 ANSWER 72 OF 89 CA COPYRIGHT 2003 ACS TI Differential block of nicotinic synapses on B versus Differential block of nicotinic synapses on B versus C neurons in sympathetic ganglia of frog by .alpha-conotoxins MII and Iml PY

MII on .alpha.3.beta.2 neuronal nicotinic receptors

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ユロ ANSWER 73 OF 89 CA COPYRIGHT 2003 ACS

Identification of genes encoding A-lineage ***conotoxin*** peptides by PCR PY 1996 1995 1996

ユロ ANSWER 74 OF 89 CA COPYRIGHT 2003 ACS

L4 ANSWER 75 OF 89 CA COPYRIGHT 2003 ACS TI Use of ***conotoxin*** peptides U002 and MII for

Use of ***conotoxin*** peptides U002 and MII for treating or detecting small-cell lung carcinoma PY 1996 1997 1996 1998 1998 1999

L4 ANSWER 76 OF 89 CA COPYRIGHT 2003 ACS
TI .alpha.- ***Conotoxin*** -lml: a competitive antagonist at .alpha.-bungarotoxin-sensitive neuronal nicotinic receptors in hippocampal neurons PY 1996

L4 ANSWER 78 OF 89 CA COPYRIGHT 2003 ACS

The 1.1 ANG. crystal structure of the neuronal acetylcholine receptor antagonist, alpha - ***conotoxin*** PnIA from Conus pennaceus PY

L4 ANSWER 79 OF 89 CA COPYRIGHT 2003 ACS A new .alpha.- ***conotoxin*** which targets .alpha.3.beta.2 nicotinic acetylcholine receptors PY ANSWER 80 OF 89 CA COPYRIGHT 2003 ACS

.alpha.- ***Conotoxin*** Imperialis I inhibits nicotine-evoked hormone release and cell proliferation in human neuroendocrine carcinoma cells

L4 ANSWER 81 OF 89 CA COPYRIGHT 2003 ACS ***Conotoxin*** peptides of Conus striatus PY 1995 1996 1995 1997 1996 2002 1998 2002

I. alpha. ***Conotoxin*** EI, A New Nicotinic Acet

.alpha.- ***Conotoxin*** El, A New Nicotinic Acetylcholine Receptor Antagonist with Novel Selectivity PY

L4 ANSWER 83 OF 89 CA COPYRIGHT 2003 ACS T1 .alpha.- ***Conotoxin*** Im I exhibits subtype-spe

and .alpha.9 receptors PY alpha. ***Conotoxin*** Im I exhibits subtype-specific nicotinic acetylcholine receptor blockade: preferential inhibition of homomeric alpha.7

ANSWER 84 OF 89 CA COPYRIGHT 2003 ACS

11 alpha. Conotoxins selectively inhibit one of the two acetylcholine binding sites of nicotinic receptors PY

L4 ANSWER 85 OF 89 CA COPYRIGHT 2003 ACS

TI Conotoxins having acetylcholine receptor binding properties and their use in receptors assays and pharmaceuticals PY 1995 1997 1996 1997 1998 1995 1995 1995

L4 ANSWER 86 OF 89 CA COPYRIGHT 2003 ACS TI New Mollusk-Specific .alpha.-Conotoxins Block Api

New Mollusk-Specific .alpha.-Conotoxins Block Aplysia Neuronal Acetylcholine Receptors PY

L4 ANSWER 87 OF 89 CA COPYRIGHT 2003 ACS

TI A nicotinic acetylcholine receptor ligand of unique specificity, .alpha. ***conotoxin*** ImI PY 1994

ANSWER 88 OF 89 CA COPYRIGHT 2003 ACS

Ti Novel .alpha.- and .omega.-conotoxins and Conus striatus venom PY

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ANSWER 89 OF 89 CA COPYRIGHT 2003 ACS

Conotoxin -related compounds PY

ANSWER 6 OF 89 CA COPYRIGHT 2003 ACS AN 137:332622 CA

Toxins 'R' Us: more pharmacological tools from nature's superstore Harvey, Alan L.

ONR, UK Strathclyde Institute for Drug Research, Dept of Physiology and Pharmacology, University of Strathclyde, Glasgow, G4

Trends in Pharmacological Sciences (2002), 23(5), 201-203 CODEN: TPHSDY; ISSN: 0165-6147

Elsevier Science Ltd. DT Journal; General Review LA English

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L4 ANSWER 48 OF 89 CA COPYRIGHT 2003 ACS AN 131:82983 CA
TI Uses of alpha- ***conotoxin*** peptides

Uses of alpha- ***conotoxin*** peptides

Olivera, Baldomero M.; Mcintosh, J. Michael; Yoshikami, Doju; Cartier, G. Edward; Luo, Siqin

₽₹ University of Utah Research Foundation, USA

SO PCT Int. Appl., 40 pp. CODEN: PIXXD2 DT Patent LA English

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9933482 A1 19990708

WO 1998-US27367 19981223

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BF, BJ, CF, CG, CI,CM, GA, GN, GW, ML, MR, NE, SN, TD, TG AU 9920917 A1 19990719 AU 1999-20917 19981; US 6265541 B1 20010724 US 1998-219446 19981223 19981223

PRAI US 1997-70153P P 19971231 US 2002022715 A1 20020221 US 2001-897465 20010703

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OS MARPAT 131:82983 WO 1998-US27367 W 19981223

FORMAT RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORDALL CITATIONS AVAILABLE IN THE RE

L4 ANSWER 16 OF 89 CA COPYRIGHT 2003 ACS

AN 136:146437 CA

function and cDNAs encoding them TI New members of the .mu.- ***conotoxin*** family for use in the treatment of disease associated with sodium channel

IN Olivera, Baldomero M.; McIntosh, J. Michael; Garrett, James E.; Watkins, Maren; Cruz, Lourdes J.; Shon, Ki-Joon; Jacobsen, Richard; Jones, Robert M.; Cartier, G. Edward; Shen, Gregory S.

DT Patent LA English FAN.CNT 1 PA University of Utah Research Foundation, USA; Cognetix, Inc. SO PCT Int. Appl., 231 pp. CODEN: PIXXD2

PATENT NO. KIND DATE APPLICATION NO. DATE

PRAI US 2000-219619P P 20000721 WO 2001-US23125 W 20010723 US 2001-264319P P US 2000-245157P P 20001103 AU 2001082945 A5 20020205 US 2001-277270P P WO 2002007678 A2 20020131 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ 20010129 20010321 AU 2001-82945 WO 2001-US23125 20010723

L4 ANSWER 25 OF 89 CA COPYRIGHT 2003 ACS AN 133:146232 CA

Alpha-conotoxins and nucleic acids encoding them

Watkins, Maren; Olivera, Baldomero M.; Hillyard, David R.; McIntosh, J. Michael; Jones, Robert M.

DT Patent LA English FAN.CNT 2 University of Utah Research Foundation, USA; Cognetix, Inc. SO PCT Int. Appl., 229 pp. CODEN: PIXXD2

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2000044776 A1 20000803 WO 2000-US1979 20000128

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1147130 A1 20011024 EP 2000-908382 20000128

PRAI US 1999-118381P P 19990129 JP 2002534996 T2 20021022 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO JP 2000-596032 20000128

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OS MARPAT 133:146232

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L4 ANSWER 26 OF 89 CA COPYRIGHT 2003 ACS AN 133:131093 CA

TI Protein and cDNA sequences of Conus .alpha.-conotoxins and the therapeutic uses thereof as neuromuscular blocking

PA University of Utah Research Foundation, USA; Cognetix, Inc. SO PCT Int. Appl., 95 pp. CODEN: PIXXD2 IN Olivera, Baldomero M.; Layer, Richard T.; Watkins, Maren; Hillyard, David R.; McIntosh, J. Michael; Jones, Robert M. DT Patent LA English FAN.CNT

PATENT NO. KIND DATE APPLICATION NO. DATE

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W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
  PRAI AU 1998-5895 A 19980914
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GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   L4 ANSWER 32 OF 89 CA COPYRIGHT 2003 ACS
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Craik, David James; Daly, Norelle Lee; Nielsen, Katherine Justine
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Preparation of cyclized ***conotoxin*** peptides
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PA University of Utah Research Foundation, USA SO PCT Int. Appl., 30 pp. CODEN: PIXXD2 DT Patent LA English FAN.CNT 1
                                                                                              IN McIntosh, J. Michael; Kulak, Jennifer M.; Yoshikami, Doju; Olivera, Baldomero M.
                                                                                                                             TI Use of alpha. ***conotoxin*** MII? to treat disorders resulting from nicotine-stimulated dopamine release
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T1 Preparation and interaction of alpha.- ***conotoxin*** peptides with neuronal nicotinic acetylcholine receptors
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Olivera, Baldomero M.; Cruz, Lourdes J.; Hillyard, David R.; Mcintosh, J. Michael; Santos, Ameurfino D.
                                                                                                                                                                                                                                                                                                                                                                                                       ***Conotoxin*** peptides
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AT 1995-901691 19941019
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PRAI US 1993-84848
WO 1994-US11927 W 19941019
             US 1993-137800 A 19931019
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                            A2 19930629
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19950607
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L4 ANSWER 78 OF 89 CA COPYRIGHT 2003 ACS AN 124:335314 CA

TI The 1.1 .ANG. crystal structure of the neuronal acetylcholine receptor antagonist, .alpha.- ***conotoxin*** PnIA from Conus

Hu, Shu-Hong; Gehrmann, John; Guddat, Luke W.; Alewood, Paul F.; Craik, David J.; Martin, Jennifer L Center Drug Design and Development, Univ. Queensland, St. Lucia, 4072, Australia

Structure (London) (1996), 4(4), 417-423 CODEN: STRUE6; ISSN: 0969-2126 PB Current Biology

Journal LA English

may be important for the specific interaction of PnIA with neuronal nAChR. residues being located at opposite ends of the mol.); a hydrophobic region and a protruding tyrosine side chain. These features characterized by distinct hydrophobic and polar surfaces; a 16. ANG, sepn. of the sole pos. and neg. charges (these two charged PnIA toxin provides a rigid framework for presentation of chem. groups that are required for activity. The structure is three-dimensional structure incorporates a .beta. turn followed by two .alpha.-helical turns. The conformation is stabilized by two disulfide bridges that form the interior of the mol., with all other side chains oriented outwards. The compact architecture of the AB The 1.1 ANG, crystal structure of synthetic PnIA was detd. by direct methods using the Shake-and-Bake program. The

L4 ANSWER 81 OF 89 CA COPYRIGHT 2003 ACS AN 124:48165 CA

Conotoxin peptides of Conus striatus

Olivera, Baldomero M.; Cruz, Lourdes J.; Hillyard, David R.; McIntosh, J. Michael; Santos, Ameurfina D.

DT Patent LA English FAN.CNT 7 University of Utah Research Foundation, USA SO PCT Int. Appl., 66 pp. CODEN: PIXXD2

PI WO 9511256 W: AU, CA, JP A1 19950427 KIND DATE APPLICATION NO. DATE WO 1994-US11927 19941019

EP 728146 AU 681216 AU 9510831 US 5514774 RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE B2 19970821 A1 19960828 A1 19950508 19960507 19960828 EP 1995-901691 19941019 US 1993-137800 19931019 AU 1995-10831 19941019

EP 728146 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE T2 19980914 B1 20020109 JP 1994-512187 19941019

PRAI US 1993-137800 A 19931019 JP 10509415 12 10509415 E 20020115 AT 1995-901691 19941019

US 1993-84848 A2 19930629

WO 1994-US11927 W 19941019

L4 ANSWER 82 OF 89 CA COPYRIGHT 2003 ACS AN 123:330553 CA

AU Martinez, Jennifer S.; Olivera, Baldomero M.; Gray, William R.; Craig, A. Grey; Groebe, Duncan R.; Abramson, Stewart N.; alpha. ***Conotoxin*** El, A New Nicotinic Acetylcholine Receptor Antagonist with Novel Selectivity

Department of Biology, University of Utah, Salt Lake City, UT, 84112, USA

Biochemistry (1995), 34(44), 14519-26 CODEN: BICHAW; ISSN: 0006-2960 PB American Chemical Society

Journal LA English

nAChRs, alpha. ***conotoxin*** El selectively binds the agonist site near the alpha./ delta. subunit interface in contrast to alpha.- ***conotoxin*** MI which selectively targets the alpha./ gamma. ligand binding site. In mammalian nAChRs alpha.-***conotoxin*** EI shows high affinity for both the .alpha./.delta. and .alpha./.gamma. subunit interfaces (with some preference previously isolated .alpha.-conotoxins. We demonstrate that this structural divergence has functional consequences. In Torpedo purified from the venom of Conus ermineus and is called .alpha. - ***conotoxin *** EI. The sequence diverges from that of amino acid peptide and is the first reported .alpha.- ***conotoxin*** from an Attantic fish-hunting cone. The peptide was AB We report the isolation and characterization of a novel nicotinic acetylcholine receptor (nAChR) ligand. The toxin is an 18

> function for characterization of nAChRs. has been verified by total chem. synthesis. Thus, alpha.- ***conotoxin*** El is a newly-available tool with unique structure and bridging between Cys4-Cys10 and Cys5-Cys18, analogous to those of previously described .alpha.-conotoxins. This sequence sequence of the peptide is: Arg-Asp-Hyp-Cys-Cys-Tyr-His-Pro-Thr-Cys-Asn-Met-Ser-Asn- Pro-Gin-Ile-Cys-NH2, with disulfide for the .alpha./.delta. site), whereas .alpha.- ***conotoxin*** MI is highly selective for the .alpha./.delta. ligand binding site. The

L4 ANSWER 85 OF 89 CA COPYRIGHT 2003 ACS AN 123:50449 CA

Conotoxins having acetylcholine receptor binding properties and their use in receptors assays and pharmaceuticals

Olivera, Baldomero M.; Rivier, Jean E. F.; Cruz, Lourdes J.; Abogadie, Fe; Hopkins, Chris E.; Dykert, John; Torres, Josep L. Salk Institute for Biological Studies, USA; University of Utah Research Foundation

SO PCT Int. Appl., 55 pp. CODEN: PIXXD2

LA English

FAN.CNT 7 PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9501436 W: AU, CA, JP, KR A1 19950112 WO 1994-US7194 19940627

US 5432155 RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE A 19950711 US 1993-84848 19930629

AU 678837 AU 9471158 CA 2165566 B2 19970612 A1 19950124 AA 19950112 AU 1994-71158 19940627 CA 1994-2165566 19940627

AU 9735197 US 5700778 EP 706566 R: AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE A1 19960417 A 19971223 A1 19971120 US 1995-458499 EP 1994-920316 19940627 AU 1997-35197 19970821

PRAI US 1993-84848 AU 699078 B2 19981119 19940627 19930629

WO 1994-US7194 W MARPAT 123:50449

solid phase peptide synthesis and tested for biol. activity. acetylcholine receptors and in pharmaceuticals (no data). Thirteen different conotoxins contg. 16-46 amino acids were prepd. by ion channels. The peptides are of such length that they can be made by chem. synthesis. The peptides may be used to analyze are useful both in vivo and in assays because they specifically target particular receptors, such as the acetylcholine receptor, and AB Substantially pure conotoxins are provided which inhibit synaptic transmissions at the neuromuscular junctions and which

L4 ANSWER 86 OF 89 CA COPYRIGHT 2003 ACS AN 121:201190 CA

TI New Mollusk-Specific .alpha . Conotoxins Block Aplysia Neuronal Acetylcholine Receptors

Fainzilber, Michael; Hasson, Arik; Oren, Ruth; Burlingame, Alma L.; Gordon, Dalia; Spira, Micha E.; Zlotkin, Eliahu

Silberman Institute of Life Sciences, Hebrew University of Jerusalem, Jerusalem, 91904, Israel

AB Two mollusk-specific neurotoxic peptides from the venom of the molluscivorous snail Conus pennaceus are described Biochemistry (1994), 33(32), 9523-9 CODEN: BICHAW; ISSN: 0006-2960 DT Journal LA English

study of structure-function relationships in ACh receptors. pos. charged residue in piscivorous venom. alpha.-conotoxins. In contrast to other alpha.-conotoxins, which are selective for the depolarizing responses evoked by brief iontophoretic application of acetylcholine. The pharmacol. and amino acid sequences of the toxins (alpha.PnIA, GCCSLPPCAANNPDYC-NH2; alpha.PnIB, GCCSLPPCALSNPDYC-NH2) enable their As such they are new probes which can be used to define subtypes of ACh receptors, and they should be useful tools in the features, the most striking of which is the presence of a single neg. charged residue in the C-terminal loop. This loop contains a induces 5-10-mV membrane depolarization, which recovers to the control level within 1-3 min in the presence of the toxin. T vertebrate skeletal muscle nicotinic ACh receptors, these Conus pennaceus toxins block neuronal ACh receptors in molluscs. classification as novel .alpha.-conotoxins. The sequences differ from those of previously described.alpha.-conotoxins in a no. of response is blocked by 1 mM hexamethonium. Concomitantly with the transient depolarization, the toxins block approx. 90% These new toxins block acetylcholine receptors (AChR) of cultured Aplysia neurons. Bath application of 0.5-1 .mu.M toxin

L4 ANSWER 88 OF 89 CA COPYRIGHT 2003 ACS AN 117:206785 CA

TI Novel alpha.- and .omega.-conotoxins and Conus striatus venom

Kristipati, Ramasharma; Ramachandran, J.; Miljanich, George; et al. AU Ramilo, Cecilia A.; Zafaralla, Glenn C.; Nadasdi, Laszlo; Hammerland, Lance G.; Yoshikami, Doju; Gray, William R.;

CS Mar. Sci. Inst., Univ. Philippines, Quezon City, 1101, Philippines SO Biochemistry (1992), 31(41), 9919-26 CODEN: BICHAW; ISSN: 0006-2960

DT Journal

conotoxin SVIB is distinct from the high-affinity .omega.- ***conotoxin*** GVIA or MVIIA site Binding competition expts, with rat brain synaptosomal membranes indicate that the high-affinity binding site for omega. previously purified from Conus venoms; only omega. ***conotoxin*** SVIB has proven to be lethal to mice upon ic injection. CKLKGQSCRKTSYDCCSGSCGRSGKC-NH2. This peptide has a different pharmacol. specificity from other omega.-conotoxins vertebrate species, it was much less effective in mammals. The third toxin, omega. ***conotoxin*** SVIB, has the sequence SVIB. .omega.-SVIA is the smallest natural .omega.- ***conotoxin*** so far characterized and has the sequence CRSSGSPCGVTSICCGRCYRGKCT.NH2. Although .omega.- ***conotoxin*** SVIA is a potent paralytic toxin in lower pos. charge, and has a free C-terminus. The other 2 paralytic peptides are Ca channel-targeted .omega.-conotoxins, SVIA and GCCCNPACGPNYGCGTSCS. In contrast to all other .alpha.-conotoxins, SII has 3 disulfide bonds (instead of two), has no net One of these, an acetylcholine receptor blocker designated .alpha. ***conotoxin*** SII, has the sequence Three neurotoxic peptides from the venom of C. striatus were purified, biochem, characterized, and chem, synthesized

ANSWER 89 OF 89 CA COPYRIGHT 2003 ACS AN 105:43332 CA ***Conotoxin*** -related compounds

IN Sakakibara, Shunpei; Nishiuchi, Yuji

Ajinomoto Co., Inc., Japan SO Jpn. Kokai Tokkyo Koho, 16 pp.

CODEN: JKXXAF

Patent

FAN.CNT 1 LA Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

PRAI JP 1984-83463 PI JP 60226899 A2 19851112 19840425 JP 1984-83463 19840425

/ Structure 1 in file .gra /

the reaction was stopped by addn. of ascorbic acid in citric acid buffer (pH 5.0) to give 11% I (U = H, W = NH2, V = Pro). AcNHCH2) was dissolved in aq. MeOH/dioxane contg. HCl and 0.1N iodine/MeOH added with vigorous agitation. After 20 min AB The title compds. I [U = Gly.Arg-, H; V = Pro, Gly, W = NH2, OH], useful as muscle relaxants and hypnotics (effectiveness comparable to that of a tubocurarine deriv.), were prepd. Thus, the protected cyclic oligopeptide amine II (Acm =

L5 ANSWER 1 OF 4 CA COPYRIGHT 2003 ACS

TI Pairwise interactions between neuronal .alpha.7 acetylcholine receptors and .alpha.conotoxin PnIB PY

L5 ANSWER 2 OF 4 CA COPYRIGHT 2003 ACS

Il Single amino acid substitutions in alpha-conotoxin PnIA shift selectivity for subtypes of the mammalian neuronal nicotinic acetylcholine

ANSWER 3 OF 4 CA COPYRIGHT 2003 ACS

Uses of alpha-conotoxin peptides PY 1999 1999 2001 2002

L5 ANSWER 4 OF 4 CA COPYRIGHT 2003 ACS
TI Accelerated chemical synthesis of peptides and small proteins PY 1999

ANSWER 2 OF 4 CA COPYRIGHT 2003 ACS

132:132466 CA

TI Single amino acid substitutions in .alpha.-conotoxin PnIA shift selectivity for subtypes of the mammalian neuronal nicotinic

Hogg, Ron C.; Miranda, Les P.; Craik, David J.; Lewis, Richard J.; Alewood, Paul F.; Adams, David J.

The Department of Physiology and Pharmacology, University of Queensland, Brisbane, 4072, Australia

Journal of Biological Chemistry (1999), 274(51), 36559-36564 CODEN: JBCHA3; ISSN: 0021-9258

American Society for Biochemistry and Molecular Biology DT Journal LA English

examd., with IC50 values of 14 and 33 nM, and a maximal redn. in current amplitude of 87% and 71%, resp. The modified PnIA and PnIB were found to cause concn. dependent inhibition of the ACh-induced current in all rat parasympathetic neurons of the role played by different nAChR subtypes in cell function and communication. In this study, the native .aipha.conotoxins AB The alpha conotoxins, a class of nicotinic acetylcholine receptor (nAChR) antagonists, are emerging as important probes

> RE ONT 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD mammalian parasympathetic neurons have been dissected with these conopeptides. subtypes, including alpha.3.beta.2 and alpha.3.beta.4. Four distinct components of the nicotinic ACh-induced current in propose that positions 10 and 11 of PnIA and PnIB influence potency and det. selectivity among .alpha.7 and other nAChR indicate that PnIA, PnIB, and the analogs, [A10L]PnIA and [N11S]PnIA, have identical backbone structures. The authors an IC50 value of 1.4 nM and 46% maximal block of the total current. The residual current was not inhibited further by .alpha.caused 91% block. [A10L]PnIA was the most potent inhibitor, reducing the ACh-induced current in apprx.80% of neurons, with bungarotoxin, but was further reduced by the .alpha.-conotoxins PnIA or PnIB, and by mecamylamine. 1H NMR studies alpha.-conotoxin [N11S]PnIA reduced the ACh-induced current with an IC50 value of 375 nM and a maximally effective conon.

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